MARKED-UP VERSION OF CLAIMS

1. (amended) Peptides of the AT_1 receptor, ~~preferably~~ comprising 5 to 30~~, preferably 5 to 10~~ amino acids as well as their variants, which can form an epitope and bind auto-antibodies occurring in preeclampsia and malign hypertension.

2. Peptides according to Claim 1, wherein they comprise SEQ ID no. 1 AFHYESQ or contain this sequence in an identical or slightly modified form.

3. (amended) Peptides according to Claim 1, wherein they ~~are~~ comprise at least one of the amino acid sequences AVHYQSN, SHFYQTR, GYYFDTN or ENTNIT or contain at least one of these sequences in an identical or slightly modified form.

4. (amended) Antibodies aimed against the epitope of the AT_1 receptor, wherein they recognise the peptides according to ~~Claims 1 to 3~~claim 1.

5. Antibodies according to Claim 4, wherein they recognise the peptides of SEQ ID no. 1 or peptides with the amino acid sequence AVHYQSN, SHFYQTR, GYYFDTN or ENTNIT.

6. (amended) ~~Use of the human AT_1 receptor, preferably of the peptides according to Claims 1 to 3,~~ A method for the production of agents for diagnostic and therapeutic purposes in diseases with a

positive antibody status, in particular preeclampsia, comprising use of the human AT_1 receptor.

7. (amended) ~~Use~~The method according to Claim 6, wherein auto-antibody binding peptides according to ~~Claims 1 to 3 are~~claim 1 is used.

8. (twice amended) The method~~Use~~ according to claim 6, wherein recombinantly produced, autoantibody binding receptor parts of the AT_1 receptor as well as of the peptides according to ~~Claims 1 to 3 are used~~claim 1 are used.

9. (twice amended) The method~~Use~~ according to Claim 6, wherein peptides according to ~~Claims 1 to 3~~claim 1 and/or molecules containing these peptides are used soluble or bound to a solid phase for direct or indirect ~~(competitive)~~ detection of antibodies in body fluids~~, in particular blood~~.

10. (twice amended) The method ~~Use~~ according to Claim 6, wherein peptides according to ~~Claims 1 to 3~~claim 1 and/or molecules containing these peptides are used bound to a solid phase for binding and elimination of the pathological, functionally active autoantibodies in body fluids, ~~in particular blood, i.e.~~ for immunoglobulin adsorption.

11. (twice amended) The method ~~Use~~ according to Claim 6, wherein the amino acid sequences and/or molecules containing these sequences are used bound to a solid phase for binding and elimination of the pathological, functionally active autoantibodies in body fluids~~, in particular blood, i.e.~~ for immunoglobulin

adsorption in combination with unspecific ~~(~~overall immunoglobulin binding ligands~~)~~.

12. (amended) Method for ~~B~~binding and elimination of the pathological, functionally active autoantibodies according to ~~Claims 4 and 5~~claim 4 in body fluids, in particular blood, by use of inspecific adsorber molecules ~~such as~~chosen from the group consisting of protein A, protein G, antihuman immunoglobulin as well as overall immunoglobulin binding ligands ~~such as~~chosen from the group consisting of ~~amino acids, in particular~~ L-tryptophane ~~or~~and peptides.

13. (amended) ~~Use of peptides at least containing at least one of the amino acid sequences according to Claims 1 to 3~~ A method for the immunisation of mammals for the purpose of obtaining polyclonal and monoclonal antibodies, comprising using peptides at least containing at least one of the amino acid sequences according to claim 1.

14. (amended) Use of ~~antibodies aimed against the amino acid sequences according to Claims 1 to 3~~ A method for immunisation of mammals for the purpose of obtaining antiidiotypical antibodies, comprising using antibodies aimed against the amino acid sequences according to claim 1.

15. (amended) Antigenic agent for detection of preeclampsia and malign hypertension, wherein it contains at least one peptide according to ~~Claims 1 to 3~~claim 1~~, preferably SEQ ID no. 1~~.

16. (amended) Immunogenic agent, wherein it contains at least one peptide according to ~~Claims 1 to 3, preferably SEQ ID no. 1~~claim 1, which induces the production of antibodies capable of recognising auto-antigens in preeclampsia or malign hypertension.

17. (amended) Test kit to determine anti- AT_1 receptor antibodies for proof of preeclampsia or malign hypertension, containing at least one peptide according to ~~Claims 1 to 3~~claim 1.

18. (amended) Method ~~to detect~~for detecting anti- AT_1 receptor antibodies in biological fluids, wherein the sample to be examined is brought into contact with at least one peptide of ~~Claims 1 to 3~~claim 1 or with a combination of these peptides with a carrier material under conditions permitting an antigen-antibody reaction and rendering proof by means of physical or chemical methods~~ known per se~~.

19. (amended) ~~Use of the peptides according to Claims 1 to 3~~ A method for production of therapeutic agents against preeclampsia or malign hypertension. Comprising using the peptides according to claim 1.